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WO 9606098A1

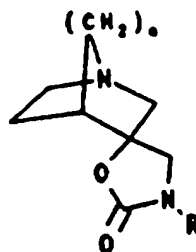
INTERNATIONAL APPLICATION PUBLISHED UNDER

(51) International Patent Classification ⁶ : C07D 498/20, 453/02		A1	(11) International Publication Number: WO 96/06098
			(43) International Publication Date: 29 February 1996 (29.02.96)
(21) International Application Number: PCT/SE95/00937		(81) Designated States: AM, AT, AU, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TT, UA, UG, US, UZ, VN, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG), ARIPO patent (KE, MW, SD, SZ, UG).	
(22) International Filing Date: 22 August 1995 (22.08.95)			
(30) Priority Data: 9417084.2 24 August 1994 (24.08.94) GB 9504627.2 8 March 1995 (08.03.95) GB			
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(54) Title: **SPIRO-AZABICYCLIC COMPOUNDS USEFUL IN THERAPY**

(57) Abstract

There are provided new compounds of formula (I), wherein R represents hydrogen or methyl; and n represents 1 or 2; or a pharmaceutically acceptable acid addition salt thereof, together with processes for preparing them, compositions containing them and their use in therapy. Compounds of formula (I) are expected to be useful in the treatment of psychotic disorders, intellectual impairment disorders and anxiety.



(I)

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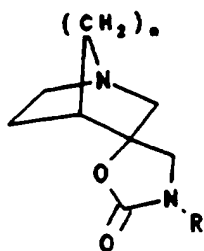
Spiro-azabicyclic compounds useful in therapy

This invention relates to novel compounds, processes for preparing them, compositions containing them and their use in therapy.

Spiro-azabicyclic compounds are known to have therapeutic activity in a range of disorders of the central nervous system. Taiwan Patent Application 201312, European Patent Application 452101 (both of Israel Institute of Biological Research), International Patent Application WO 95/03303 (Israel Institute of Biological Research; published after the earliest priority date of this application) and European Patent Application 350118 (Merck Sharpe and Dohme) disclose azabicyclic compounds including azabicyclo(2.2.2)octane and/or azabicyclo(2.2.1)heptane derivatives spiro-connected to 5-membered rings which have muscarinic agonist activity and which are indicated for the treatment of diseases caused by deficiency in central cholinergic function. European Patent Application 337547 (Merck, Sharpe and Dohme) discloses azabicyclic compounds spiro-connected to 5-membered rings which are antagonists of 5-HT₃ receptors and which are indicated for the treatment of inter alia schizophrenia, nausea, migraine and Alzheimer's disease.

We have now identified a new group of spiro-azabicyclic compounds which have useful pharmacological properties.

Thus, according to the invention, we provide a compound of formula I:



wherein

R represents hydrogen or methyl; and

n represents 1 or 2;

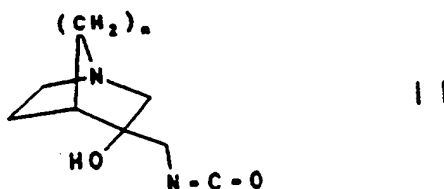
and pharmaceutically acceptable acid addition salts thereof.

We prefer compounds of formula I in which R represents hydrogen. We prefer compounds of formula I in which n represents 2.

- 5 As a second aspect of the invention we provide a process for the preparation of a compound of formula I, or a pharmaceutically acceptable acid addition salt thereof, which comprises:

- (a) preparing a compound of formula I in which R represents hydrogen by cyclising a corresponding compound of formula II

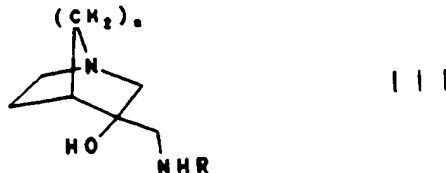
10



- 15 wherein n is as defined above;

- (b) preparing a compound of formula I by reacting a corresponding compound of formula III

20



wherein n and R are as defined above
with a carbonyl donating compound; or

- 25 (c) preparing a compound of formula I in which R represents methyl by alkylating a corresponding compound of formula I in which R represents hydrogen;
(d) preparing one enantiomer of a compound of formula I by resolving the one enantiomer from a mixture of enantiomers;

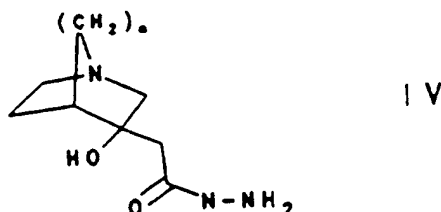
and where desired or necessary converting the resultant compound of
30 formula I, or an acid addition salt thereof, to a pharmaceutically acceptable acid addition salt thereof, or vice versa.

In process (a) the reaction will take place on warming the compound of formula II in a polar protic solvent e.g. water.

In process (b), examples of carbonyl donating compounds include carbonyldiimidazole, carbonyldichloride (phosgene) and triphosgene. The ring closure reaction will take place on heating or refluxing the compound of formula III with carbonyldiimidazole in a polar organic solvent such as THF for 1-4 hours or until reaction is complete. Alternatively phosgene can be bubbled through a solution of the compound of formula III in an organic solvent such as THF or toluene at elevated temperature for 1-4 hours or until reaction is complete.

In process (c), the alkylation reaction will take place under conditions well known in the art, for example, by treating the compound of formula I in which R represents hydrogen with a strong base followed by a methyl halide e.g. methyl iodide.

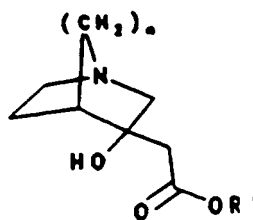
Compounds of formula II may be prepared by Curtius rearrangement of a compound of formula IV:



wherein n is as defined above.

Typical reaction conditions for the Curtius rearrangement are discussed in J March "Advanced Organic Chemistry" (1985) 3rd Edition, pages 984-5 however we prefer to perform the reaction by treating the compound of formula IV in water with sodium nitrite and warming to approximately 85°C for approximately 1 hour. The compound of formula IV is not isolated and the cyclisation reaction described in process (a) proceeds directly in situ.

Compounds of formula IV may be prepared by treatment of a compound of formula V

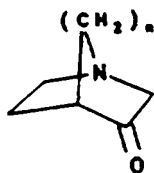


V

wherein n is as defined above, and R' is an alkyl or aryl group,
with anhydrous hydrazine.

This reaction may be performed in a polar protic solvent at ambient temperature over 4-12 hours. We prefer that R' represents alkyl, typically methyl, ethyl or t-butyl.

Compounds of formula V may be prepared by reaction of a compound of formula VI



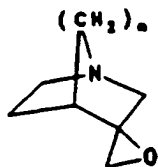
VI

wherein n is as defined above
with an acetic acid ester in the presence of n-butyl lithium. The acetic acid ester (for example the t-butyl or ethyl ester) is first treated with n-butyl lithium at -78 °C and the reaction proceeds on adding the compound of formula VI and warming to room temperature. The product is obtained on quenching with water.

Compounds of formula VI are either known or may be prepared by known methods.

Compounds of formula III may be prepared by reacting a compound of formula

VII



5 wherein n is as defined above,
with ammonia or methylamine.

This reaction may be performed under standard conditions for example by combining the reagents in a polar protic solvent at ambient or elevated temperature.

10 Compounds of formula III may also be prepared by reacting a compound of formula VI with trimethylsilylcyanide (Me_3SiCN) followed by reduction with, for example, lithium aluminium hydride or Raney Nickel. Reaction conditions will be familiar to a person skilled in the art.

15 Compounds of formula VII may be prepared from the corresponding compounds of formula VI using one of the reagents well known in the art for preparation of oxiranes from ketones (see, for example, the reactions referenced in J March "Advanced Organic Chemistry" (1985) 3rd Edition, page 1161). We have found that trimethylsulphoxonium iodide is a suitable reagent and the reaction may be performed
20 in DMSO at a temperature of between room temperature and 80 °C over a period of up to 2 hours, or until reaction is complete.

Acid addition salts of the compounds of formula I which may be mentioned include salts of mineral acids, for example the hydrochloride and hydrobromide salts; and salts
25 formed with organic acids such as formate, acetate, malate, benzoate and fumarate salts.

Acid addition salts of compounds of formula I may be formed by reacting the free base or a salt, enantiomer or protected derivative thereof, with one or more equivalents of the appropriate acid. The reaction may be carried out in a solvent or medium in which
30 the salt is insoluble or in a solvent in which the salt is soluble, eg water, dioxan, ethanol, tetrahydrofuran or diethyl ether, or a mixture of solvents, which may be removed in vacuo or by freeze drying. The reaction may be a metathetical process or it may be carried out on an ion exchange resin.

The compounds of the invention and intermediates may be isolated from their reaction mixtures by standard techniques.

5 The compounds of formula I exist in tautomeric or enantiomeric forms, all of which are included within the scope of the invention. The various optical isomers may be isolated by separation of a racemic mixture of the compounds using conventional techniques, e.g. fractional crystallisation, or chiral HPLC. Alternatively the individual enantiomers may be made by reaction of the appropriate optically active starting materials under reaction conditions which will not cause racemisation.

10 Intermediate compounds also exist in enantiomeric forms and may be used as purified enantiomers, racemates or mixtures.

The compounds of the invention are indicated as pharmaceuticals, in particular in the
15 treatment or prophylaxis of psychotic disorders and intellectual impairment disorders. Examples of psychotic disorders include schizophrenia, mania and manic depression. Examples of intellectual impairment disorders include Alzheimer's disease, learning deficit, cognition deficit and attention deficit hyperactivity disorder. The compounds of the invention may also be useful as analgesics and in the treatment or prophylaxis of
20 Parkinson's and Huntington's disease, neurodegenerative disorders in which there is loss of cholinergic synapses, Tourette's syndrome, depression and anxiety. The compounds may further be indicated for the treatment or prophylaxis of jetlag and for use in inducing the cessation of smoking.

25 According to a further aspect of the invention we provide a compound of the invention for use as a pharmaceutical, especially in the treatment or prophylaxis of the aforementioned diseases or conditions.

For the above-mentioned uses the dosage administered will, of course, vary with the
30 compound employed, the mode of administration and the treatment desired. However, in general, satisfactory results are obtained when the compounds of the invention are administered at a daily dosage of from about 0.1mg to about 20mg per kg of animal body weight, preferably given in divided doses 1 to 4 times a day or in sustained release

form. For man, the total daily dose is in the range of from 5mg to 1,400mg, more preferably from 10mg to 100mg, and unit dosage forms suitable for oral administration comprise from 2mg to 1,400mg of the compound admixed with a solid or liquid pharmaceutical carrier or diluent.

5 The compounds of formula I, and pharmaceutically acceptable salts thereof, may be used on their own or in the form of appropriate medicinal preparations for enteral or parenteral administration. According to a further aspect of the invention, there is provided a pharmaceutical formulation including preferably less than 80% and more
10 preferably less than 50% by weight of a compound of the invention in admixture with a pharmaceutically acceptable diluent or carrier.

Examples of diluents and carriers are:

for tablets and dragees: lactose, starch, talc, stearic acid;

15 for capsules: tartaric acid or lactose;

for injectable solutions: water, alcohols, glycerin, vegetable oils;

for suppositories: natural or hardened oils or waxes.

There is also provided a process for the preparation of such a pharmaceutical composition
20 tion which comprises mixing the ingredients.

According to a further aspect of the invention, there is provided the use of a compound of formula I, or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for the treatment or prophylaxis of one of the above mentioned diseases
25 or conditions; and a method of treatment or prophylaxis of one of the above mentioned diseases or conditions, which comprises administering a therapeutically effective amount of a compound of formula I, or a pharmaceutically acceptable salt thereof, to a patient.

Compounds of formula IV and compounds of formula III in which R represents methyl
30 are new and useful intermediates. Thus a further aspect of the invention we provide a compound of formula IV, or a salt thereof. We also provide compounds of formula III in which R represents methyl, or a salt thereof.

Compounds of formula I are agonists of nicotinic acetylcholine receptors.

While not being limited by theory, it is believed that agonists of the $\alpha 7$ nAChR (nicotinic acetylcholine receptor) subtype should be useful in the treatment or prophylaxis of psychotic disorders and intellectual impairment disorders, and have advantages over compounds which are or are also agonists of the $\alpha 4$ nAChR subtype. Therefore, compounds which are selective for the $\alpha 7$ nAChR subtype are preferred.

The pharmacological activity of the compounds of the invention may be measured in the tests set out below.

Test A

Assay for affinity at $\alpha 7$ nAChR subtype

125 I- α -Bungarotoxin (BTX) binding to rat hippocampal membranes

Rat hippocampi were homogenized in 20 volumes of cold homogenization buffer (HB: concentrations of constituents (mM): tris(hydroxymethyl)aminomethane 50; MgCl_2 1; NaCl 120; KCl 5; pH 7.4). The homogenate was centrifuged for 5 minutes at 1000g, the supernatant was saved and the pellet re-extracted. The pooled supernatants were centrifuged for 20 minutes at 12000g, washed, and resuspended in HB. Membranes (30-80 μg) were incubated with 5 nM [125 I] α -BTX, 1 mg/ml BSA (bovine serum albumin), test drug, and either 2 mM CaCl_2 or 0.5 mM EGTA [ethylene glycol-bis(β -aminoethyl-ether)] for 2 hours at 21°C, and then filtered and washed 4 times over Whatman glass fibre filters (thickness C) using a Brandel cell harvester. Pretreating the filters for 3 hours with 1% BSA/0.01% PEI (polyethyleneimine) in water was critical for low filter blanks (0.07 % of total counts per minute). Nonspecific binding was described by 100 μM (-)-nicotine, and specific binding was typically 75%.

Test B

Assay for affinity to the $\alpha 4$ nAChR subtype

[^3H]-(-)-nicotine binding

Using a procedure modified from Martino-Barrows and Kellar (Mol Pharm 31:169-174; 1987), rat brain (cortex and hippocampus) was homogenized as in the [125 I] α -BTX binding assay, centrifuged for 20 minutes at 12000g, washed twice, and then resuspended in HB containing 100 μM diisopropyl fluorophosphate. After 20 minutes at 4°C, mem-

branes (≈ 0.5 mg) were incubated with 3 nM [^3H]-(-)-nicotine, test drug, 1 μM atropine, and either 2 mM CaCl_2 or 0.5 mM EGTA for 1 hours at 4°C , and then filtered over Whatman glass fibre filters (thickness C) (pretreated for 1 hour with 0.5% PEI) using a Brandel cell harvester. Nonspecific binding was described by 100 μM carbachol, and specific binding was typically 84%.

Binding data analysis for Tests A and B

IC₅₀ values and pseudo Hill coefficients (n_H) were calculated using the non-linear curve fitting program ALLFIT (DeLean A, Munson P J and Rodbard D (1977) Am. J. Physiol., 235:E97-E102). Saturation curves were fitted to a one site model, using the non-linear regression program ENZFITTER (Leatherbarrow, R.J. (1987)), yielding K_D values of 1.67 and 1.70 nM for the ^{125}I - α -BTX and [^3H]-(-)-nicotine ligands respectively. K_i values were estimated using the general Cheng-Prusoff equation:

$$K_i = [\text{IC}_{50}] / ((2 + ([\text{ligand}] / [K_D])^n)^{1/n} - 1)$$

where a value of $n=1$ was used whenever $n_H < 1.5$ and a value of $n=2$ was used when $n_H \geq 1.5$. Samples were assayed in triplicate and were typically $\pm 5\%$. K_i values were determined using 6 or more drug concentrations.

When compared with compounds of the prior art, the compounds of the invention have the advantage that they may be less toxic, be more efficacious, be longer acting, have a broader range of activity, be more potent, produce fewer side effects, are more easily absorbed or have other useful pharmacological properties.

The invention is illustrated by the following examples:

Example 1

Spiro[1-azabicyclo[2.2.2]octane-3,5'-oxazolidine]-2'-one monohydrochloride

(a) 2-(3-Hydroxy-1-azabicyclo[2.2.2]oct-3-yl)acetic acid 'butyl ester

To a 0°C solution of diisopropylamine (6.7 ml) in tetrahydrofuran (THF) (20 ml) was added 2.3M $n\text{BuLi}$ (20 ml). The reaction mixture was stirred for 40 minutes and then

cooled to -78°C . To this mixture was added dropwise a solution of 'butylacetate (6.4 ml) in 10 ml of THF and stirring continued for an additional 15 minutes. Quinuclidin-3-one (free base) (5.0 g) in THF (15 ml) was added to the mixture dropwise and the mixture was allowed to warm to 0°C over 1 hour. To this solution was added water (100 ml), the solution was extracted twice with chloroform and the combined extracts were washed once with brine. The resulting solution was dried over MgSO_4 , filtered and dried in vacuo to give 9.53 g of the subtitle compound as an off-white solid.

(b) 2-(3-Hydroxy-1-azabicyclo[2.2.2]oct-3-yl)acetic acid hydrazide

To a solution of 3.5 g of the compound of step (a) in 15 ml of methylene chloride was added trifluoroacetic acid (39 ml) and the mixture was stirred at ambient temperature for three hours. The mixture was then concentrated in vacuo. The residue was dissolved in methanol (30 ml) and 18M H_2SO_4 (3 ml) was added and the mixture was stirred overnight. The mixture was then poured into a solution of sodium carbonate in water, extracted three times with chloroform, dried over MgSO_4 , filtered and concentrated in vacuo to give a yellow solid. The solid was dissolved in methanol (10 ml) and hydrazine (2 ml) was added to this solution and the mixture was heated to reflux for one hour. The mixture was then concentrated in vacuo. The resulting solid was suspended in toluene (50 ml) and heated to reflux in an apparatus equipped with a Dean-Stark trap for two hours. The reaction mixture was then allowed to cool, and the product was filtered off to give 1.82 g of the subtitle compound as a tan solid.

(c) Spiro[1-azabicyclo[2.2.2]octane-3,5'-oxazolidine]-2'-one monohydrochloride

To a solution of 0.91 g of the compound of step (b) in water (7 ml) was added 12M HCl to give a solution with a pH of 1. The mixture was cooled to 0°C and a 0°C solution of sodium nitrite (0.33 g) in 5 ml of water was added. The solution was stirred at 0°C for twenty minutes and then heated to 70°C for an additional twenty minutes. The reaction mixture was cooled to ambient temperature and basified with 50% $\text{NaOH}/\text{H}_2\text{O}$. The solution was saturated with NaCl and extracted with chloroform 4x20 ml, dried over MgSO_4 , filtered and concentrated in vacuo. The residue was dissolved in methanol and HCl gas was bubbled through the mixture until the pH was less than 2. Diethyl ether was added to the solution and the resulting white solid was filtered off to give 0.73 g of the title compound as an off-white solid, m.p. $289-291^{\circ}\text{C}$. ^1H NMR (500 MHz, DMSO): 1.7-1.9 (m, 3H), 2.05 (m, 1H), 2.25 (bs, 1H), 3.15 (m, 3H), 3.2-3.35 (m, 1H), 3.4-3.6 (m, 3H), 3.6 (d, 1H), 7.75 (s, 1H), 10.95 (bs, 1H).

Example 2

(+)- and (-)-Spiro[1-azabicyclo[2.2.2]octane-3,5'-oxazolidine]-2'-one monohydrochloride

To a solution of the compound of Example 1 (3.8 g) in absolute ethanol was added a solution of dibenzoyl-L-tartaric acid (7.474 g) in absolute ethanol and to this mixture was added a small portion of ethyl acetate. A solid formed upon standing within an hour which was filtered off and the filtrate retained. The solid was dissolved in aqueous sodium hydroxide, extracted with chloroform 3x50 ml, dried over MgSO_4 , filtered and concentrated in vacuo. The residue was dissolved in methanol and HCl gas was bubbled through the mixture until the pH was less than 2. Diethyl ether was added to the solution and the resulting white solid was filtered off to give 0.63 g of the (+)-enantiomer, m.p. $>250^\circ\text{C}$. $[\alpha]_D = +57.978$ ($c = 0.6570$, CH_3OH).

The filtrate retained above was converted to the free base by concentrating the solution and then dissolving the residue in 20% $\text{NaOH}/\text{H}_2\text{O}$, which was extracted with chloroform, dried over MgSO_4 , filtered and concentrated in vacuo to give 1.21 g of residue. The residue was dissolved in absolute ethanol and combined with a solution of 2.38 g of dibenzoyl-D-tartaric acid in absolute ethanol. The mixture was stirred and a small portion of ethyl acetate added resulting in the formation of a white solid after stirring for an hour. The precipitate was filtered and dissolved in 20% $\text{NaOH}/\text{H}_2\text{O}$, extracted with chloroform, dried over MgSO_4 , filtered and concentrated in vacuo. The residue was dissolved in methanol and HCl gas was bubbled through the mixture until the pH was less than 2. Diethyl ether was added to the solution and the resulting white solid was filtered off to give 0.21 g of the (-)-enantiomer, m.p. $>250^\circ\text{C}$. $[\alpha]_D = -61.473$ ($c = 0.7727$, CH_3OH).

Example 3

Spiro[1-azabicyclo[2.2.1]heptan-3,5'-oxazolidin-2'-one] monohydrochloride

(a) 3-Hydroxy-1-azabicyclo[2.2.1]hept-3-yl acetic acid ethyl ester

To a cooled (-78°C) solution of diisopropylamine (2.65 ml, 0.0203 moles) in tetrahydrofuran (150 ml) was added 2.5 M n-BuLi in hexanes (8.1 ml, 0.0203 moles) dropwise over one minute. After ten minutes, ethyl acetate (1.97 ml, 0.0203 moles) was added dropwise over one minute. After a further ten minutes, 1-azabicyclo[2.2.1]heptan-3-one (1.5 g, 0.0135 moles, prepared by the method of J Chem Soc. Chem Commun, 1618, 1988)

in tetrahydrofuran (25 ml) was added dropwise over five minutes. After twenty five minutes, the cold bath was removed and the mixture was allowed to warm to ambient temperature. To this mixture was added water (100 ml) followed by chloroform (250 ml). The organic layer was separated and the aqueous layer was extracted with chloroform (100 ml). The combined organic extracts were dried over MgSO_4 , filtered and concentrated *in vacuo* to give the subtitle compound (1.12 g).

(b) 3-Hydroxy-1-azabicyclo[2.2.1]hept-3-yl acetic acid hydrazide

To a solution of the product of step (a) (1.12 g, 0.0056 moles) in methanol (20 ml) at room temperature was added anhydrous hydrazine (1.76 ml, 0.056 moles) and the mixture was stirred overnight. The mixture was concentrated *in vacuo* and azeotroped twice with toluene (100 ml) by heating to reflux in an apparatus fitted with a Dean-Stark trap to give the subtitle compound (1.11 g).

(c) Spiro[1-azabicyclo[2.2.1]heptan-3,5'-oxazolidin-2'-one] monohydrochloride

To a cooled (0°C) solution of the product of step (b) (1.11 g, 0.006 moles) in water (50 ml) adjusted to pH 1 with concentrated HCl was added dropwise over two minutes an aqueous solution of NaNO_2 (0.455 g, 0.0066 moles). After the addition was complete the cold bath was removed and the mixture was heated to 85°C for one hour. The mixture was allowed to cool to ambient temperature and basified to pH 10 with 50% $\text{NaOH}/\text{H}_2\text{O}$. This mixture was extracted three times with chloroform (150 ml), dried over MgSO_4 , filtered and concentrated *in vacuo*. The residue was flash chromatographed on silica gel with methanol/ethyl acetate [1:1]. The appropriate fractions were collected and concentrated. The residue was then dissolved in chloroform, filtered and concentrated *in vacuo*. The residue was dissolved in ethyl acetate/methanol and treated with a saturated solution of HCl/ethyl acetate until the solution had a pH < 6. The solution was concentrated to give 120 mg of residue. This residue was dissolved in hot methanol and precipitated with ether to give the title compound (28 mg), m.p. $>250^\circ\text{C}$. ^1H NMR (500 MHz, DMSO): 1.95 (m, 1H), 2.1 (m, 1H), 3.0 (s, 1H), 3.2-3.4 (m, 5H), 3.5-3.6 (m, 3H), 7.8 (s, 1H), 10.6 (bs, 1H).

Example 43'-Methyl spiro-[1-azabicyclo[2.2.2]octan-3,5'-oxazolidin-2'-one] monohydrochloride(a) Spiro-1-azabicyclo[2.2.2]octan-3-oxirane

To a suspension of sodium hydride (11.6 g of 60% oil dispersion washed 3x with hexane: 0.35 mol) in 600 mL of DMSO was added 101.4 g (0.35 mol) of trimethylsulfoxonium iodide in portions. After gas evolution had subsided, the mixture was stirred at room temperature for 30 min. A solution of quinuclidin-3-one (44 g, 0.35 mol) in 200 mL of THF was added dropwise over a period of 30 min and the mixture was warmed to 70-80 °C for 1 h. The reaction mixture was cooled to RT, poured onto 1 L of ice water and the aqueous solution extracted with CHCl₃ (4 x 300 mL). The organic extracts were washed with H₂O (3 x 500 mL), saturated brine (1 x 300 mL), dried over MgSO₄ and the solution evaporated to afford the subtitle compound as a yellow oil (37.6 g, 77%).

(b) 3-Hydroxy-1-azabicyclo[2.2.2]oct-3-yl methylaminomethane

To a solution of 25 mL (0.72 mol) of condensed methylamine in 75 mL of methanol was added 10.2 g (0.073 mol) of the product of step (a) and the solution stirred at room temperature for 16 h. The reaction mixture was concentrated under reduced pressure and the resulting oil was dissolved in 75 mL of CHCl₃ and concentrated to afford 12.4 g (100%) of the subtitle compound as an oil.

(c) 3'-Methyl spiro-[1-azabicyclo[2.2.2]octan-3,5'-oxazolidin-2'-one]monohydrochloride

To a solution of the product of step (b) (8.0 g, 0.047 mol) in 100 mL of dry THF was added 9.15 g (0.056 mol) of carbonyldiimidazole and the mixture was heated at reflux for 3 h. The reaction mixture was then cooled to room temperature and concentrated under vacuum. The residual oil was dissolved in 200 mL of CH₂Cl₂, the solution washed with H₂O (1 x 100 mL), brine (1 x 50 mL), dried over MgSO₄ and evaporated to afford a yellow solid. The solid was dissolved in 50 mL of 1:1 methanol/isopropanol mixture, acidified with HCl in methanol, and the precipitate collected, washed with cold methanol and dried to afford 5.6 g (51%) of the title compound as a colorless solid, m.p. 305-307 °C. ¹H NMR (500 MHz, DMSO): 1.8 (m, 3H), 2.05 (m, 1H), 2.25 (bs, 1H), 2.75 (s, 3H), 3.1-3.2 (m, 3H), 3.25-3.35 (m, 1H), 3.4-3.5 (m, 1H), 3.5-3.6 (m, 2H), 3.7 (d, 1H), 10.9 (bs, 1H).

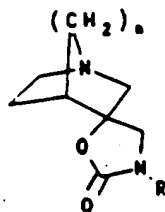
Example 5

- The compounds spiro[1-azabicyclo[2.2.2]octane-3,5'-oxazolidine]-2'-one monohydrochloride, (+)-spiro[1-azabicyclo[2.2.2]octane-3,5'-oxazolidine]-2'-one monohydrochloride, (-)-spiro[1-azabicyclo[2.2.2]octane-3,5'-oxazolidine]-2'-one monohydrochloride, spiro[1-azabicyclo[2.2.1]heptan-3,5'-oxazolidin-2'-one] monohydrochloride and 3'-methyl spiro-1-azabicyclo[2.2.2]octan-3,5'-oxazolidin-2'-one monohydrochloride were tested in Test A above and gave binding affinities (K_i) of less than $10\mu\text{M}$ indicating that they are expected to have useful therapeutic activity.
- 10 The compounds spiro[1-azabicyclo[2.2.2]octane-3,5'-oxazolidine]-2'-one monohydrochloride, (+)-spiro[1-azabicyclo[2.2.2]octane-3,5'-oxazolidine]-2'-one monohydrochloride, (-)-spiro[1-azabicyclo[2.2.2]octane-3,5'-oxazolidine]-2'-one monohydrochloride, spiro[1-azabicyclo[2.2.1]heptan-3,5'-oxazolidin-2'-one] monohydrochloride and 3'-methyl spiro-1-azabicyclo[2.2.2]octan-3,5'-oxazolidin-2'-one monohydrochloride were tested in Test B
15 above and gave K_i values at least 1.6 times greater than those given in Test A indicating that they possess desirable selectivity.

Claims:

15

1. A compound of formula I:



wherein

R represents hydrogen or methyl; and

- 10 n represents 1 or 2:

or a pharmaceutically acceptable acid addition salt or enantiomer thereof.

2. A compound according to claim 1 wherein R represents hydrogen, or a pharmaceutically acceptable acid addition salt or enantiomer thereof.

3. Spiro[1-azabicyclo[2.2.2]octane-3,5'-oxazolidine]-2'-one, or a pharmaceutically acceptable acid addition salt or enantiomer thereof.

4. (+)-Spiro[1-azabicyclo[2.2.2]octane-3,5'-oxazolidine]-2'-one, or a pharmaceutically acceptable acid addition salt thereof.

5. (-)-Spiro[1-azabicyclo[2.2.2]octane-3,5'-oxazolidine]-2'-one, or a pharmaceutically acceptable acid addition salt thereof.

6. Spiro[1-azabicyclo[2.2.1]heptan-3,5'-oxazolidin-2'-one], or a pharmaceutically acceptable acid addition salt or enantiomer thereof.

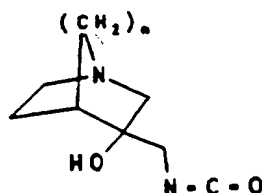
7. 3'-Methyl spiro-[1-azabicyclo[2.2.2]octan-3,5'-oxazolidin-2'-one], or a pharmaceutically acceptable acid addition salt or enantiomer thereof.

8. A pharmaceutical formulation including a compound as defined in any one of claims 1 to 7, or a pharmaceutically acceptable acid addition salt or enantiomer thereof, in admixture with a pharmaceutically acceptable diluent or carrier.

9. A compound according to any one of claims 1 to 7, or a pharmaceutically acceptable acid addition salt or enantiomer thereof, for use as a pharmaceutical.

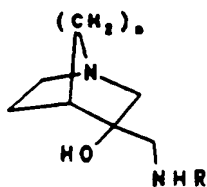
10. The use of a compound as defined in any one of claims 1 to 7, or a pharmaceutically acceptable acid addition salt or enantiomer thereof, in the manufacture of a medicament for the treatment or prophylaxis of psychotic disorders or intellectual impairment disorders.

11. The use as claimed in claim 10, wherein the disorder is schizophrenia, Alzheimer's disease, learning deficit, cognition deficit or attention deficit hyperactivity disorder.
12. The use of a compound as defined in any one of claims 1 to 7, or a pharmaceutically acceptable acid addition salt or enantiomer thereof, in the manufacture of a medicament for the treatment or prophylaxis of anxiety.
13. A method of treatment or prophylaxis of psychotic disorders or intellectual impairment disorders, which comprises administering a therapeutically effective amount of a compound as defined in any one of claims 1 to 7, or a pharmaceutically acceptable acid addition salt or enantiomer thereof, to a patient.
14. The method as claimed in claim 13, wherein the disorder is schizophrenia, Alzheimer's disease, learning deficit, cognition deficit or attention deficit hyperactivity disorder.
15. A method of treatment or prophylaxis of anxiety, which comprises administering a therapeutically effective amount of a compound as defined in any one of claims 1 to 7, or a pharmaceutically acceptable acid addition salt or enantiomer thereof, to a patient.
16. A process for preparation of a compound of formula I, as defined in claim 1, or a pharmaceutically acceptable acid addition salt or enantiomer thereof, which comprises:
- (a) preparing a compound of formula I in which R represents hydrogen by cyclising a corresponding compound of formula II



wherein n is as defined in claim 1:

- (b) preparing a compound of formula I by reacting a corresponding compound of formula III



I I I

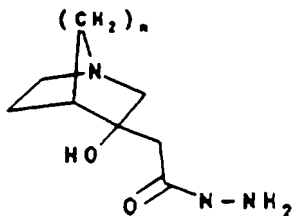
wherein n and R are as defined in claim 1,
with a carbonyl donating compound;

(c) preparing a compound of formula I in which R represents methyl by alkylating
a corresponding compound of formula I in which R represents hydrogen; or

(d) preparing one enantiomer of a compound of formula I by resolving the one en-
antiomer from a mixture of enantiomers;

and where desired or necessary converting the resultant compound of
formula I, or an acid addition salt or enantiomer thereof, to a pharmaceutically accept-
able acid addition salt or enantiomer thereof, or vice versa.

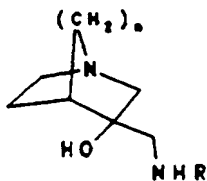
17. A compound of formula IV



I V

wherein n is as defined in claim 1, or a salt or enantiomer thereof.

18. A compound of formula III



I I I

wherein n is as defined in claim 1 and R represents methyl, or a salt or enantiomer
thereof.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 95/00937

A. CLASSIFICATION OF SUBJECT MATTER

IPC: C07D 498/20, C07D 453/02

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC: C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

CA, WPI

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,A	WO 9503303 A2 (KOSMIN, GERALD, EMMANUEL), 2 February 1995 (02.02.95) --	1-12,16-18
A	EP 0452101 A2 (ISRAEL INSTITUTE FOR BIOLOGICAL RESEARCH), 16 October 1991 (16.10.91) --	1-12,16-18
A	WO 9201690 A1 (FISONS CORPORATION), 6 February 1992 (06.02.92) --	1-12,16-18

☒ Further documents are listed in the continuation of Box C.☒ See patent family annex.

- * Special categories of cited documents:
- * "A" document defining the general state of the art which is not considered to be of particular relevance
- * "E" earlier document but published on or after the international filing date
- * "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- * "O" document referring to an oral disclosure, use, exhibition or other means
- * "P" document published prior to the international filing date but later than the priority date claimed
- * "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- * "X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- * "Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
- * "&" document member of the same patent family

Date of the actual completion of the international search

28 November 1995

Date of mailing of the international search report

28.11.95

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INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 95/00937

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	STN International, File CA. Chemical Abstracts, volume 80, no. 23, 10 June 1974, (Columbus Ohio, US), Nakanishi, Michio et al: "Quinuclidine derivatives", abstract no. 133284, & JP B4 49007159 740219 Showa, JP 70-128283 701225 --	1-12
A	STN International, File CA, Chemical abstracts, volume 78, no. 11, 19 March 1973, Columbus, Ohio, US), Maillard, Jacques et al: "Cycloalkanespiro-heterocyclic compounds. VIII. S-4'-Piperidino) spirooxazolidin-2-ones (I-oxa 3,8-diazaä4, 5äspirodecan-2-ones)", abstract no. 71968, & Chim. Ther. (1972), 7(6), 458-66 --	1-12
A	DE 2505143 A1 (BUSKINE S.A.), 14 August 1975 (14.08.75) -- -----	17

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 95/00937

Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: 13-15
because they relate to subject matter not required to be searched by this Authority, namely:
See PCT Rule 39.1(iv): Methods for treatment of the human or animal body by surgery or therapy, as well as diagnostic methods.
2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT
Information on patent family members

30/10/95

International application No.
PCT/SE 95/00937

Patent document cited in search report		Publication date	Patent family member(s)		Publication date
WO-A2-	9503303	02/02/95	NONE		
EP-A2-	0452101	16/10/91	IL-A-	97726	29/12/94
			JP-A-	6293761	21/10/94
			US-A-	5053412	01/10/91
			US-A-	5407938	18/04/95
WO-A1-	9201690	06/02/92	EP-A-	0540625	12/05/93
			US-A-	5073560	17/12/91
DE-A1-	2505143	14/08/75	AU-B,B-	498871	29/03/79
			AU-A-	7806975	12/08/76
			BE-A,A-	825444	29/05/75
			CA-A-	1033363	20/06/78
			CH-A-	601304	14/07/78
			FR-A,A-	2262666	26/09/75
			GB-A-	1495647	21/12/77
			JP-A-	50112376	03/09/75
			US-A-	4028351	07/06/77